Prisca 5.1.0.17

Date of report: 16-11-2024

NA

Patient data			
Name	Mrs. KUNTI DEVI	Patient ID	0522411130066
Birthday	01-02-2004	Sample ID	a1357729
Age at sample date	20.8	Sample Date	13-11-2024
Gestational age	11 + 4		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	49	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

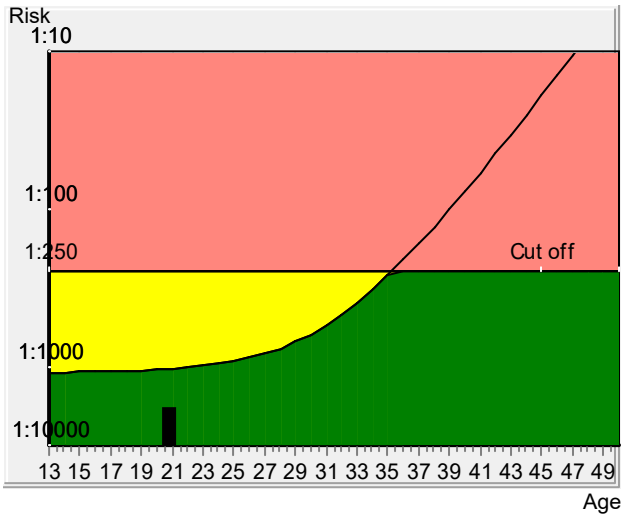
Parameter	Value	Corr. MoM
PAPP-A	2.33 mIU/mL	0.64
fb-hCG	65.03 ng/ml	1.25

Risks at sampling date

Age risk	1:1040
Biochemical T21 risk	1:1396
Combined trisomy 21 risk	1:7264
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	11 + 0
Method	CRL Robinson
Scan date	09-11-2024
Crown rump length in mm	44
Nuchal translucency MoM	0.90
Nasal bone	present
Sonographer	NA
Qualifications in measuring NT	MD



Trisomy 21

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
After the result of the Trisomy 21 test (with NT) it is expected that among 7264 women with the same data, there is one woman with a trisomy 21 pregnancy and 7263 women with not affected pregnancies.
The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.
Please note that risk calculations are statistical approaches and have no diagnostic value!
The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!

Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

Sign of Physician

below cut off | Below Cut Off, but above Age Risk | above cut off